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U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

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TRANSMITTAL LETTER TO THE UNITED STATES
 DESIGNATED/ELECTED OFFICE (DO/EO/US)
 CONCERNING A FILING UNDER 35 U.S.C. 371

RUH-284

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

09/937524

INTERNATIONAL APPLICATION NO.
 PCT/EP00/02455

INTERNATIONAL FILING DATE
 March 21, 2000

PRIORITY DATE CLAIMED
 March 27, 1999

TITLE OF INVENTION CATALYSTS FOR THE GAS-PHASE OXIDATION OF ETHYLENE AND
 ACETIC ACID TO VINYL ACETATE, A PROCESS FOR PRODUCING THEM AND THEIR USE

APPLICANT(S) FOR DO/EO/US
 HAGEMEYER et al

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☒ This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.
4. ☐ The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☒ is attached hereto (required only if not communicated by the International Bureau).
 - b. ☐ has been communicated by the International Bureau.
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☒ An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
 - a. ☒ is attached hereto.
 - b. ☐ has been previously submitted under 35 U.S.C. 154(d)(4).
7. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are attached hereto (required only if not communicated by the International Bureau).
 - b. ☐ have been communicated by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has **NOT** expired.
 - d. ☐ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☒ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☒ An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11 to 20 below concern document(s) or information included:

11. ☒ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
 Three
12. ☒ Assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☒ A **FIRST** preliminary amendment.
14. ☐ A **SECOND** or **SUBSEQUENT** preliminary amendment.
15. ☐ A substitute specification.
16. ☐ A change of power of attorney and/or address letter.
17. ☐ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
18. ☐ A second copy of the published international application under 35 U.S.C. 154(d)(4).
19. ☐ A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
20. ☒ Other items or information: PCT/RO/101

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21. ☒ The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)):

Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.443(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO. \$1000.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$860.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.443(a)(2)) paid to USPTO \$710.00

International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$690.00

International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00

ENTER APPROPRIATE BASIC FEE AMOUNT =

CALCULATIONS PTO USE ONLY

\$1000.00

\$ 1000.00

Surcharge of \$130.00 for furnishing the oath or declaration later than months from the earliest claimed priority date (37 CFR 1.492(e)). ☐ 20 ☐ 30

CLAIMS **NUMBER FILED** **NUMBER EXTRA** **RATE**

Total claims - 20 = x \$18.00

Independent claims - 3 = x \$80.00

MULTIPLE DEPENDENT CLAIM(S) (if applicable) + \$270.00

TOTAL OF ABOVE CALCULATIONS = \$ 1000.00

☐ Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.

SUBTOTAL = \$ 1000.00

Processing fee of \$130.00 for furnishing the English translation later than months from the earliest claimed priority date (37 CFR 1.492(f)). ☐ 20 ☐ 30

TOTAL NATIONAL FEE = \$ 1000.00

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property + 120.00

TOTAL FEES ENCLOSED = \$ 1120.00

Amount to be refunded: \$

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a. ☐ A check in the amount of \$ _____ to cover the above fees is enclosed.

b. ☐ Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.

c. ☒ The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 02-2275 A duplicate copy of this sheet is enclosed.

d. ☒ Fees are to be charged to a credit card. **WARNING:** Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.

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20311

PATENT, TRADEMARK OFFICE

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NAME

19,683

REGISTRATION NUMBER

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Our Ref.: RUH-284

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of: :
PCT/EP00/02455 : PCT Date: March 21, 2000
HAGEMEYER et al :
Serial No.: :
Filed: Concurrently Herewith :
For: CATALYSTS...AND THEIR USE :
600 Third Avenue
New York, NY 10016
Dated: September 24, 2001

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, D.C. 20231

Sir:

Please amend this application as follows:

IN THE SPECIFICATION:

Page 1, before line 1, insert

--This application is a 371 of PCT/EP00/02455 filed March
21, 2000.--

IN THE CLAIMS:

Claim 1 (amended) A catalyst comprising palladium, at least
one alkali metal compound and, optionally, at least one promoter on
a porous support, obtained by loading the porous support, which
comprises a reducible metal oxide of elements of groups IIIB, IVB,
VB, VIB of the Periodic Table of the Elements or ZnO or comprises

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a mixture of these oxides or a mixed oxide of these elements in which zinc may also be present, with at least one palladium compound, subsequently carrying out reduction at a temperature of 300-600°C and additionally applying at least one alkali metal compound and, optionally, at least one promoter before or after the reduction.

Claim 2 (amended) A catalyst of claim 1 which comprises at least one potassium compound.

Claim 3 (amended) A catalyst of claim 1 which additionally comprises at least one member of the group consisting of Au, Ba, Cd and their compounds as promoter.

Claim 4 (amended) A catalyst of claim 1, wherein the reducible support is TiO_2 .

Claim 5 (amended) A catalyst of claim 1, wherein the reduction is carried out for 1 minute to 24 hours.

Claim 6 (amended) A catalyst of claim 1, wherein the reduction is carried out using gaseous or vaporizable reducing agents.

Claim 7 (amended) A catalyst of claim 1, wherein the reducing agent for the reduction is at least one member selected

from the group consisting of H_2 , CO, ethylene, NH_3 , formaldehyde, methanol, hydrocarbons and mixtures of these reducing agents with inert gases.

Claim 8 (amended) A process for producing a catalyst of claim 1, comprising loading the porous support, which comprises a reducible metal oxide of elements of groups IIIb, IVb, V, VIb of the Periodic Table of the Elements or ZnO or comprises a mixture of these oxides or a mixed oxide of these elements in which zinc may also be present, with at least one palladium compound, subsequently carrying out reduction at a temperature of 300-600°C and additionally applying at least one alkali metal compound and, optionally, at least one promoter before or after the reduction.

Claim 9 (amended) The process of claim 8, wherein the catalyst comprises at least one potassium compound.

Claim 10 (amended) The process of claim 8 wherein the catalyst additionally comprises at least one member of the group consisting of Au, Ba, Cd and their compounds as promoters.

Claim 11 (amended) The process of claim 8, wherein the reducible support is TiO_2 .

Claim 12 (amended) The process of claim 8, wherein the reduction is carried out from 1 minute to 24 hours.

Claim 13 (amended) The process of claim 8, wherein the reduction is carried out using gaseous or vaporizable reducing agents.

Claim 14 (amended) The process of claim 8, wherein the reducing agent for the reduction is at least one member selected from the group consisting of H₂, CO, ethylene, NH₃, formaldehyde, methanol, hydrocarbons and mixtures of these reducing agents with inert gases.

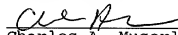
Cancel claim 15 and add the following claim.

--16. In a process for the preparation of vinyl acetate from the gaseous phase reaction of ethylene, acetic acid and oxygen or oxygen containing gas, the improvement comprising using as catalyst the catalyst of claim 1.--

REMARKS

This amendment is submitted to insert reference to the PCT application and to conform the claims to the American practice.

Respectfully submitted,
BIERMAN, MUSERLIAN AND LUCAS


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CAM:sd

Enclosures: Marked-up Version of Claims
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MARKED-UP VERSION OF CLAIMS

Claim 1 (amended) A catalyst comprising palladium, at least one alkali metal compound and, [if desired] optionally, at least one [or more promoters] promoter on a porous support, [obtainable] obtained by loading the porous support, which comprises a reducible metal oxide of elements of groups IIIB, IVB, VB, VIB of the Periodic Table of the Elements or ZnO or comprises a mixture of these oxides or a mixed oxide of these elements in which zinc may also be present, with at least one palladium compound, subsequently carrying out [a] reduction at a temperature of 300-600°C and additionally applying at least one alkali metal compound and, [if desired] optionally, at least one [or more promoters] promoter before or after the reduction.

Claim 2 (amended) A catalyst [as claimed in] of claim 1 which comprises at least one potassium compound.

Claim 3 (amended) A catalyst [as claimed in] of claim 1 [or 2] which additionally comprises at least one member of the group consisting of Au, Ba, [and/or] Cd and[/or] their compounds as [promoters] promoter.

Claim 4 (amended) A catalyst [as claimed in one or more] of [claims 1 to 3] claim 1, wherein the reducible support is TiO₂.

Claim 5 (amended) A catalyst [as claimed in one or more] of [claims 1 to 4] claim 1, wherein the reduction is carried out for [a time in the range from] 1 minute to 24 hours.

Claim 6 (amended) A catalyst [as claimed in one or more] of [claims 1 to 5] claim 1, wherein the reduction is carried out using gaseous or vaporizable reducing agents.

Claim 7 (amended) A catalyst [as claimed in one or more] of [claims 1 to 6] claim 1, wherein the reducing agent for the reduction is at least one member selected from the group consisting of H₂, CO, ethylene, NH₃, formaldehyde, methanol, hydrocarbons [and their mixtures] and mixtures of these reducing agents with inert gases.

Claim 8 (amended) A process for producing [catalysts, which comprises] a catalyst of claim 1, comprising loading the porous support, which comprises a reducible metal oxide of elements of groups IIIB, IVB, V, VIB of the Periodic Table of the Elements or ZnO or comprises a mixture of these oxides or a mixed oxide of these elements in which zinc may also be present, with at least one palladium compound, subsequently carrying out [a] reduction at a temperature of 300-600°C and additionally applying at least one alkali metal compound and, [if desired] optionally, at least one [or more promoters] promoter before or after the reduction.

Claim 9 (amended) The process [as claimed in] of claim 8, wherein the catalyst comprises at least one potassium compound.

Claim 10 (amended) The process [as claimed in] of claim 8 [or 9] wherein the catalyst additionally comprises at least one member of the group consisting of Au, Ba, [and/or] Cd and[/or] their compounds as promoters.

Claim 11 (amended) The process [as claimed in one or more] of [claims 8 to 10] claim 8, wherein the reducible support is TiO_2 .

Claim 12 (amended) The process [as claimed in one or more] of [claims 8 to 11] claim 8, wherein the reduction is carried out [for a time in the range] from 1 minute to 24 hours.

Claim 13 (amended) The process [as claimed in one or more] of [claims 8 to 12] claim 8, wherein the reduction is carried out using gaseous or vaporizable reducing agents.

Claim 14 (amended) The process [as claimed in one or more] of [claims 8 to 13] claim 8, wherein the reducing agent for the reduction is at least one member selected from the group consisting of H_2 , CO, ethylene, NH_3 , formaldehyde, methanol, hydrocarbons [and their mixtures] and mixtures of these reducing agents with inert gases.

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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As printed		
(54) Title:	CATALYSTS FOR THE GAS-PHASE OXIDATION OF ETHYLENE AND ACETIC ACID TO VINYL ACETATE AND METHOD FOR THE PRODUCTION AND USE THEREOF	
(54) Bezeichnung:	KATALYSATOREN FÜR DIE GASPHASENOXIDATION VON ETHYLEN UND ESSIGSÄURE ZU VINYLACETAT, VERFAHREN ZU IHRER HERSTELLUNG UND IHRE VERWENDUNG	
(57) Abstract		
<p>The invention relates to a supported catalyst and the production and use thereof in the synthesis of vinyl acetate in the gaseous phase from ethylene, acetic acid and oxygen, or from gases containing oxygen, whereby the activity and selectivity are simultaneously improved. The catalyst contains palladium, at least one alkali metal compound and optionally one or more catalyst accelerators, in particular, gold, barium and/or cadmium on a porous support. Said support contains at least one reducible metal oxide, in particular oxides of the elements of groups IIIB, IVB, VB, VIB from the periodic table of elements. Once the support has been loaded with at least one palladium compound, a reduction is carried out at a temperature of > 200 °C.</p>		
(57) Zusammenfassung		
<p>Die vorliegende Erfindung betrifft einen Trägerkatalysator, dessen Herstellung sowie dessen Verwendung bei der Synthese von Vinylacetat in der Gasphase aus Ethylen, Essigsäure und Sauerstoff oder Sauerstoff enthaltenden Gasen bei gleichzeitiger Verbesserung der Aktivität und Selektivität. Der Katalysator enthält Palladium, mindestens eine Alkalimetallverbindung sowie gegebenenfalls einen oder mehrere Promotoren, insbesondere Gold, Barium und/oder Cadmium auf einem porösen Träger, der mindestens ein reduzierbares Metalloxid, insbesondere Oxide der Elemente der Gruppen IIIB, IVB, VB, VIB des Periodensystems der Elemente enthält, und wobei man nach der Beladung des Trägers mit mindestens einer Palladiumverbindung eine Reduktion bei einer Temperatur von > 200 °C durchführt.</p>		

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JCO9 Rec'd PCT/PTO 25 SEP 2001
PCI/EP00/02455

Catalysts for the gas-phase oxidation of ethylene and
acetic acid to vinyl acetate, a process for producing
them and their use

- 5 The present invention relates to highly selective catalysts for the synthesis of vinyl acetate from ethylene and acetic acid, to a process for producing them and to their use.
- 10 The preparation of vinyl acetate (VAM) in the gas phase from ethylene, acetic acid and oxygen or oxygen-containing gases is known; the supported catalysts used for this synthesis comprise Pd as active metal and an alkali metal as promoter, preferably K in the form of
- 15 the acetate. Further additives used are Cd, Au or Ba.

According to US-A-4 902 823, US-A-3 939 199, US-A-4 668 819 the catalytically active components are applied in finely divided form to the catalyst support

20 by impregnation, spraying on, vapor deposition, dipping or precipitation. The procedure described gives catalysts in which the active components have penetrated to the core of the support.

- 25 However, catalysts in which the active components have not penetrated to the core but are present only in a more or less thick outer part of the support particles, i.e. the shell of the support particles, are also known (EP-A-0 565 952, EP-A-0 634 214, EP-A-0 634 209,
- 30 EP-A-0 634 208).

The catalysts known from the prior art comprise, as support, the known inert support materials such as silica, aluminum oxide, aluminosilicates, silicates,

35 titanium oxide, silicon carbide and carbon and are produced by subjecting the impregnated catalyst precursors to a reduction, e.g. in the gas phase at temperatures of 150-170°C, for example using ethylene or H₂, or in the liquid phase at temperatures of <

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100°C, e.g. using hydrazine. Higher reduction temperatures are deliberately avoided since appreciable sintering of the noble metal particles to form larger agglomerates having a lower catalytic activity occurs at temperatures > 200°C.

However, it is also known that the supports impregnated with the catalytically active components can be exposed to higher temperature ranges during the course of the production of the finished catalysts. Thus, US-A-5 336 802 and US-A-5 194 417 describe the treatment of palladium- and gold-containing catalysts by single and multiple sequential oxidation and reduction steps.

EP-A-0 839 793 relates to a process for preparing vinyl acetate in the presence of a palladium-containing catalyst during whose production the impregnated catalyst precursor which has been treated with a reducing agent is subjected to an additional sintering step at a temperature in the range from 500 to 1000°C.

WO 98/18553 discloses calcining a palladium- and gold-containing catalyst after the impregnation step at a temperature of from 100 to 600°C in a nonreducing atmosphere and only then carrying out the reduction step. The calcination can also be carried out in an oxygen atmosphere.

In the processes disclosed in EP-A-0 839 793 and WO 98/18553, the preparation of vinyl acetate is also carried out using supported catalysts based on inert support materials such as silica, aluminum oxide, aluminosilicates, silicates, titanium oxide, silicon carbide and carbon.

In view of the fact that catalysts are used in processes carried out on an industrial scale, for example the preparation of vinyl acetate, there is

great interest in improving the catalysts, particularly in respect of their activity and selectivity.

It is therefore an object of the present invention to
5 provide catalysts for the synthesis of vinyl acetate, which catalysts have a particularly high activity, selectivity and operating life.

A further object of the present invention is to provide
10 catalysts which have a particularly high stability. In particular, the catalysts should be insensitive to local temperature fluctuations and hot spots and also to fluctuations in the oxygen concentration during the reaction in the synthesis of vinyl acetate and to
15 mechanical stress.

These objects are achieved by a catalyst comprising palladium, at least one alkali metal compound and, if desired, one or more promoters on a porous support,
20 which catalyst is obtainable by loading the porous support in which at least one reducible metal oxide is present with at least one palladium compound, subsequently carrying out a reduction at a temperature of $> 200^{\circ}\text{C}$ and additionally applying at least one
25 alkali metal compound and, if desired, one or more promoters before or after the reduction.

The invention accordingly provides a catalyst comprising palladium, at least one alkali metal
30 compound and, if desired, one or more promoters on a porous support and obtainable by loading the porous support in which at least one reducible metal oxide is present with at least one palladium compound, subsequently carrying out a reduction at a temperature
35 of $> 200^{\circ}\text{C}$ and additionally applying at least one alkali metal compound and, if desired, one or more promoters before or after the reduction.

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The invention further provides a process for producing catalysts, which comprises loading the porous support in which at least one reducible metal oxide is present with at least one palladium compound, subsequently
5 carrying out a reduction at a temperature of $> 200^{\circ}\text{C}$ and additionally applying at least one alkali metal compound and, if desired, one or more promoters before or after the reduction.

10 The invention further provides a process for preparing vinyl acetate in the gas phase from ethylene, acetic acid and oxygen or oxygen-containing gases in the presence of a catalyst comprising palladium, at least one alkali metal compound and, if desired, one or more
15 promoters on a porous support and obtainable by loading the porous support in which at least one reducible metal oxide is present with at least one palladium compound, subsequently carrying out a reduction at a temperature of $> 200^{\circ}\text{C}$ and additionally applying at
20 least one alkali metal compound and, if desired, one or more promoters before or after the reduction.

In the following, the reduction of the porous support in which at least one reducible metal oxide is present
25 and which is laden with palladium compounds at a temperature of $> 200^{\circ}\text{C}$ is referred to as the high-temperature reduction and abbreviated as HTR.

An essential aspect of the catalysts of the invention
30 is the use of a support material in which at least one reducible metal oxide is present, for example oxides of the elements of groups IIIB, IVB, VB, VIB of the Periodic Table of the Elements, ZnO and other reducible oxides known from the literature and also mixtures of
35 these reducible metal oxides. Furthermore, the mixed oxides derived from the abovementioned oxides are also suitable. Preference is given to TiO_2 and ZrO_2 having intermediate to high specific surface areas as support material.

However, it is also possible to use inert support materials such as silica, aluminum oxide, aluminum silicates, silicates, silicon carbide or carbon as long as these inert materials are first coated with the
5 abovementioned reducible metal oxides. Such support materials are known from EP-A-723 810. The reducible metal oxide content is from 0.1 to 25% by weight, based on the total support material.

10 It is likewise possible to use mixtures of the reducible metal oxides with these inert materials as supports for the catalysts of the invention. Here, the proportion of the inert materials can be up to 75% by weight, based on the total support material. The
15 proportion of the inert materials is preferably up to 50% by weight, based on the total support material.

Support materials in which at least one reducible metal oxide is present are hereinafter referred to as
20 reducible support materials.

Surprisingly, nonreducible support materials known from the prior art, for example silica or aluminum oxide or mixtures thereof, are unsuitable for producing the
25 catalysts of the invention. Oxides which can~~not~~ be reduced to the metal under the production conditions employed according to the invention are likewise unsuitable as support materials.

30 The reducible support can be used either in powder form or as shaped bodies. The support material is preferably used as shaped bodies and is in the form of pellets, spheres, tablets, rings, extrudates, ribbed extrudates, star extrudates, stars, hollow extrudates or other
35 industrial shaped bodies. The diameter or the length and thickness of the support materials is generally from 3 to 9 mm. The surface area of the support measured by the BET method is generally 10-500 m²/g, preferably 15-250 m²/g. The pore volume is generally

from 0.2 to 1.2 ml/g. The pore volume is determined by means of mercury porosimetry.

5 The support materials employed according to the invention can be produced by any method known to those skilled in the art. They are generally commercially available.

10 Loading a reducible support with at least one palladium compound and subsequently carrying out a high-temperature reduction at temperatures of $> 200^{\circ}\text{C}$ and additionally applying at least one alkali metal compound and, if desired, one or more promoters before
15 or after the reduction gives catalysts which have an excellent activity and selectivity and also excellent adhesion of noble metal to the support.

20 The novel measure of the HTR of porous reducible supports laden with palladium compounds is carried out at temperatures of $> 200^{\circ}\text{C}$, preferably $> 300^{\circ}\text{C}$.

25 If desired, one or more promoters are subsequently applied to the support which has been loaded in this way. Preference is given to using at least one Au, Ba or Cd compound or a mixture of these compounds.

30 However, the addition of promoters is preferably carried out before the HTR treatment, either in admixture with the palladium compound or compounds or separately therefrom. The order plays no role; the porous reducible support can be loaded first with the palladium compound and then with the promoters or vice versa.

35 The catalysts of the invention further comprise at least one alkali metal compound, preferably at least one potassium compound. The addition of alkali metal to the porous reducible support can be carried out prior to the HTR treatment, either separately or in admixture

with the other promoters and/or with the palladium compound or compounds, or after the HTR treatment, either separately or in admixture with the other promoters.

Finally, the catalysts of the invention can be finished in accordance with the prior art and used in industrial processes, for example in the synthesis of vinyl acetate.

Γ loaded with

According to the invention, the reducible support material is first a palladium compound. The type of compound is not critical here, as long as high dispersion of the metal can be achieved. Soluble palladium compounds are suitable for this purpose, in particular water-soluble salts. Preference is given to, for example, palladium compounds selected from the group consisting of palladium(II) acetate, palladium(II) chloride, sodium tetrachloropalladate(II) $[\text{Na}_2\text{PdCl}_4]$ and palladium(II) nitrate. Besides palladium(II) acetate, further carboxylates of palladium can be used, preferably the salts of aliphatic monocarboxylic acids having from 3 to 5 carbon atoms, for example the propionate or butyrate.

However, in the case of the chlorides, it has to be ensured that the chloride ions are removed to leave a tolerable residual amount before use of the catalyst. For this purpose, the catalyst support is washed with water after loading with palladium compounds and with the promoters and after the HTR. This is achieved by washing the support, e.g. with water, after Pd and, if used, Au have been fixed to the support by reduction to the metals.

The palladium compounds mentioned can be prepared by any method known to those skilled in the art. However, most of them are also commercially available.

Promoters used are compounds of the elements Au, Ba and Cd, in particular soluble salts of these elements.

- Suitable gold compounds include, for example,
- 5 tetrachloroauric(III) acid $[\text{HAuCl}_4]$, gold(III) acetate $[\text{Au}(\text{OAc})_3]$, potassium aurate $[\text{KAuO}_2]$. It is advisable to prepare the gold acetate or the potassium aurate freshly by precipitation of the oxide/hydroxide from solutions of tetrachloroauric acid, washing and
- 10 isolating the precipitate and taking it up in acetic acid or KOH. The gold compounds are preferably applied to the support before the high-temperature reduction.

- Suitable cadmium compounds include, for example,
- 15 cadmium acetate $\text{Cd}(\text{OAc})_2$ and other carboxylates of cadmium, for example the propionate or butyrate. The cadmium compounds can be applied to the support either before or after the high-temperature reduction.

- 20 Suitable barium compounds include, for example, barium acetate $\text{Ba}(\text{OAc})_2$, other barium carboxylates such as the propionate or butyrate and also barium hydroxide $\text{Ba}(\text{OH})_2$. The barium compounds can be applied to the support either before or after the high-temperature
- 25 reduction.

- The catalyst of the invention further comprises at least one alkali metal compound, preferably at least one potassium or rubidium compound and particularly
- 30 preferably at least one potassium compound. Suitable potassium compounds include, for example, potassium acetate KOAc, potassium carbonate K_2CO_3 , potassium hydrogen carbonate KHCO_3 and potassium hydroxide KOH, and also all potassium compounds which are converted
- 35 into the acetate under the reaction conditions. The potassium compounds can be applied to the support either before or after the high-temperature reduction.

Preference is given to using the acetates for loading the porous reducible support since these catalysts cause very little chloride contamination.

- 5 The metal compounds are usually used in concentrations of from about 0.1 to 100 g per liter, preferably from 1 to 50 g per liter, based on the solvent.

- 10 Suitable solvents are all compounds in which the compounds or salts chosen are soluble and which can be easily removed again by drying after the impregnation. Examples of suitable solvents for the acetates are, in particular, unsubstituted carboxylic acids, especially acetic acid, while the chlorides are soluble primarily
15 in water or dilute hydrochloric acid.

- If the salts are not sufficiently soluble in acetic acid or in water, the additional use of a further solvent in addition to water or acetic acid, which are
20 also used as a mixture, can be advantageous.

- Possible additional solvents are ones which are inert and miscible with acetic acid or water. Examples of additions to acetic acid are ketones such as acetone
25 and acetylacetone, also ethers such as tetrahydrofuran or dioxane, acetonitrile, dimethylformamide and also hydrocarbons such as benzene.

- The application of the active component Pd and the
30 further promoters and also the alkali metal compound can be carried out by the methods known from the prior art.

- The finished catalysts have the following metal
35 contents, expressed in gram of metal per 1 l of finished catalyst:

Palladium:	generally	1 - 20 g/l
	preferably	3 - 15 g/l
	in particular	5 - 10 g/l

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reducible support and this interaction effects an advantageous change in the properties of the noble metal particles for catalysis and is responsible for excellent mechanical anchoring to the support and thus
5 a high agglomeration resistance.

The high-temperature-reduced catalysts of the invention have a particularly uniform Pd/Au active metal distribution and a high degree of dispersion of the
10 noble metals.

The high degree of dispersion is largely maintained even in long-term operation owing to reduced agglomeration of the noble metal particles, as a result
15 of which the deactivation of the catalysts of the invention is slowed and long operating lives are achieved.

In addition, the catalysts of the invention have excellent mechanical stabilities since the chemically reactive reducible supports can easily be shaped to
20 produce mechanically hard shaped bodies and the HTR at high temperatures further increases the hardness of the shaped bodies.

The catalysts of the invention are insensitive to hot spots and also to fluctuations in the oxygen concentration during the reaction of the vinyl acetate synthesis, and the intensity of the hot spots is
25 greatly reduced owing to the extraordinarily high selectivity (drastic reduction in the strongly exothermic total oxidation to CO or CO₂), as a result of which regulation and process control are significantly simplified.
30

Since an increase in the space-time yield (STY) accompanies a selectivity increase, particularly high throughputs can be realized in large plants.
35

Achievement of the abovementioned advantages could not have been foreseen from the prior art, since the prior art puts forward the opinion that reduction temperatures of $> 200^{\circ}\text{C}$ are damaging and, owing to agglomeration of the noble metal particles, reduce the activity. Thus, hot spots of $> 190^{\circ}\text{C}$ are avoided if possible both in the conventional reduction and later in operation.

10 Furthermore, the prior art teaches the use of inert supports such as silicon dioxide or silicon dioxide/aluminum oxide mixtures, preferably silicon dioxide.

15 Loading of the support with the desired amounts of the respective compounds can be carried out in one step or a plurality of sequential steps, with drying steps being able, if desired, to be inserted between the individual steps.

20 It is possible, for example, to apply one compound, e.g. a salt, of each of the elements to be applied to the support particles, e.g. Pd/K/Au, Pd/K/Cd or Pd/K/Ba. However, a plurality of salts of one element
25 can also be applied or mixed compounds of the various metals can be used. In general, exactly one salt of each of the three elements is used.

The salts can be applied to the support by known
30 loading methods, for example steeping, impregnation, spraying on, vapor deposition, dipping or precipitation. The support can be impregnated right through with the salts or it is possible to use all methods known to those skilled in the art which are
35 employed for producing surface-impregnated catalysts. These are disclosed, for example, in the documents
DE-A-1 668 088, US-A 3 775 342, US-A 3 822 308,
US-A 4 048 096, US-A 5 185 308, US-A 5 567 839,

US-A 5 314 858, EP-A-0 634 208, EP-A-0 634 209 or
EP-A-0 634 214.

In the case of the Pd/Au/K catalysts, it has been found
to be advantageous to apply the two noble metals in the
form of a shell on the support, i.e. the noble metals
are distributed only in a zone close to the surface
while the regions further inside the support body are
virtually free of noble metals. The thickness of this
catalytically active shell is usually 0.1-2 mm.

In many cases, surface-impregnated catalysts make
possible a more selective process than is the case for
catalysts in which the support particles are
impregnated to the core ("fully impregnated").

If the reaction conditions when using surface-
impregnated catalysts are left unchanged from the
reaction conditions when using fully impregnated
catalysts, more vinyl acetate can be produced per unit
reactor volume and time, which equates to a capacity
increase without additional capital costs. The work-up
of the crude vinyl acetate obtained is also made
easier, since the vinyl acetate content of the product
gas from the reactor is higher, which also leads to an
energy saving in the work-up. Suitable work-up
processes are described, for example, in
US-A 5 066 365, DE-34 22 575, DE-A-34 08 239,
DE-A-29 45 913, DE-A-26 10 624, US-A 3 840 590.

If, on the other hand, the plant capacity is kept
constant, the reaction temperature can be lowered and
the reaction can therefore be carried out more
selectively at the same total output, thus saving
starting materials. This also reduces the amount of
carbon dioxide which is formed as by-product and
therefore has to be discharged and decreases the loss
of entrained ethylene associated with this discharge.

Furthermore, this operating procedure leads to a lengthening of the operating life of the catalyst.

- According to the invention, the catalyst has to be subjected to a high-temperature reduction after loading with at least one palladium compound.

- For this purpose, a vaporizable or gaseous reducing agent can be passed at temperatures of above 200°C, preferably in the range from 200°C to 700°C and particularly preferably in the range from 300°C to 600°C, over the loaded catalyst.

- Suitable reducing agents for the HTR are all materials which are able to reduce the palladium compounds and, if used, gold compounds to the metals at the high reduction temperatures employed according to the invention.

- Preference is given to gaseous or vaporizable reducing agents for example H₂, CO, ethylene, NH₃, formaldehyde, methanol and hydrocarbons in general and also mixtures of these reducing agents. Hydrogen is particularly preferred.

- The gaseous reducing agents can also be diluted with inert gas, for example carbon dioxide, nitrogen or argon. Preference is given to using a reducing agent diluted with inert gas. Mixtures of hydrogen with nitrogen or argon, preferably those having a hydrogen content of from 1% by volume to 15% by volume, are preferred.

- The reduction times are generally in the range from 1 minute to 24 hours, particularly preferably from 30 minutes to 10 hours.

The amount of reducing agent is selected so that at least the number of equivalents of reducing agent

- required for complete reduction of the noble metals are passed over the catalyst during the treatment time. Preference is given to passing an excess of reducing agent over the catalyst in order to ensure complete
- 5 reduction. The volume flow of reducing gas is calculated from the pressure employed in the reduction, the dilution and the reaction time. Preference is given to carrying out the reduction at atmospheric pressure, i.e. at an absolute pressure of about 1 bar. For the
- 10 production of industrial amounts of the catalyst of the invention, preference is given to using a rotary tube furnace or a fluidized-bed reactor in order to ensure uniform reduction of the catalyst.
- 15 In the reduction according to the invention, it is first and foremost the noble metal compounds, i.e. Pd and for example Au, which are reduced to the corresponding metals, with the support material being partially reduced, e.g. to form Ti^{3+} centers in the TiO_2
- 20 lattice. In contrast thereto, the other metal compounds present, i.e. the alkali metal compounds and the other promoters apart from gold, are generally not reduced. For this reason, the alkali metal compounds and the nonreducible promoters can be applied to the support
- 25 either before or after the high-temperature reduction.

- In a preferred embodiment of the invention, TiO_2 or ZrO_2 supports or mixtures thereof or mixtures with further inert support constituents, for example SiO_2 and/or
- 30 Al_2O_3 , preferably as shaped bodies, more preferably in the form of pellets, spheres, rings, extrudates or tablets, are impregnated with palladium acetate and gold acetate, dried and reacted with gaseous reducing agents, preferably H_2 , ethylene and/or NH_3 , in the HTR
- 35 in the temperature range of 200-700°C, preferably 300-600°C, for a period ranging from 1 minute to 10 hours. If desired, this is followed by impregnation with potassium acetate and final drying at a temperature of

not more than 150°C, preferably 80-150°C and in particular 100-150°C.

However, it is also possible to use support materials in which the inert support constituents have firstly been coated with TiO_2 or ZrO_2 or with a mixture of these oxides. This is followed by the impregnation step and the HTR. The proportion of TiO_2 or ZrO_2 is from 0.1 to 25% by weight, based on the support material.

Further pretreatment or after-treatment steps which are known to those skilled in the art can be inserted. These include, inter alia, washing, drying, calcination, oxidation and/or reduction.

The catalysts of the invention are preferably used for preparing vinyl acetate. This is generally carried out by passing acetic acid, ethylene and oxygen or oxygen-containing gases at temperatures of from 100 to 220°C, preferably from 120 to 200°C, and pressures of from 1 to 25 bar, preferably from 1 to 20 bar, over the finished catalyst, with unreacted components being able to be circulated. The oxygen concentration is advantageously kept below 10% by volume (based on the gas mixture excluding acetic acid). However, dilution with inert gases such as nitrogen or carbon dioxide is sometimes also advantageous. Carbon dioxide is particularly suitable for dilution, since it is formed in small amounts during the reaction. The vinyl acetate formed is isolated by means of suitable methods as are described, for example, in US-A 5 066 365, DE-A-34 22 575, DE-A-34 08 239, DE-A-29 45 913, DE-A-26 10 624, US-A 3 840 590.

The following examples serve to explain and illustrate the invention without restricting it thereto.

Example 1

- 2.11 g of palladium acetate (224.49 g/mol) and 1.32 g of gold acetate (374.10 g/mol) were dissolved in 30 ml of glacial acetic acid. The preparation of gold acetate is described, for example, in US 4,933,204. 100 ml of TiO₂ supports (P25 pellets, DEGUSSA) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at 70°C on a rotary evaporator, residues of solvent were then removed at 60°C with the aid of an oil pump and finally in a vacuum drying oven for a period of 14 hours, likewise at 60°C.
- The reduction was carried out using a gas mixture of 10% by volume of H₂ in N₂. Here, the gas was passed at a temperature of about 500°C through the pellets for one hour. The reduction was carried out under atmospheric pressure at a flow of 40 l/h of a mixture of 10% by volume of H₂ in nitrogen. To load them with potassium ions, the pellets were treated with a solution of 4 g of potassium acetate in 30 ml of water. This mixture was allowed to act on the pellets at room temperature for 15 minutes in a mixer. The solvent was subsequently removed on a rotary evaporator. The pellets were dried for 14 hours at 110°C in a drying oven.

The catalyst comprised: 7 g/l Au; 16 g/l K; <1 g/l Cl; 10 g/l Pd.

- The production of the catalyst was repeated twice. The testing of these catalysts in the synthesis of vinyl acetate is reported as Examples 1A to 1C in Table 1 below.

Example 2

2.11 g of palladium acetate (224.49 g/mol) and 1.32 g of gold acetate (374.10 g/mol) were dissolved in 45 ml

of glacial acetic acid. 100 ml of TiO_2 supports (XT25376 pellets, Norton) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at 70°C on a rotary evaporator, residues of solvent were then removed at 60°C with the aid of an oil pump and finally in a vacuum drying oven for a period of 14 hours, likewise at 60°C .

- 10 The reduction was carried out using a gas mixture consisting of 10% by volume of H_2 in N_2 . Here, the gas was passed at a temperature of about 500°C through the pellets for 1 hour. The reduction was carried out under atmospheric pressure at a flow of 40 l/h of a mixture
- 15 of 10% by volume of H_2 in nitrogen. To load them with potassium ions, the pellets were treated with a solution of 4 g of potassium acetate in 45 ml of water. The pellets which had been impregnated in this way were mixed for 10 minutes at room temperature. The solvent
- 20 was subsequently removed on a rotary evaporator. The pellets were dried for 14 hours at 110°C in a drying oven.

The catalyst comprised: 7 g/l Au; 16 g/l K; <1.1 g/l Cl; 10 g/l Pd.

Examples 3A-C

- 30 2.11 g of palladium acetate (224.49 g/mol), 1.32 g of gold acetate (374.10 g/mol) and 4.0 g of potassium acetate were dissolved in 30 ml of glacial acetic acid. 100 ml of TiO_2 supports (P25 pellets, DEGUSSA) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at
- 35 60°C on a rotary evaporator and residues of solvent were then removed at 60°C in a vacuum drying oven over a period of 4 hours.

The batch was divided into three parts which were subjected to different reduction conditions.

Batch	Time	Temperature	Reducing agent
A	4 h	400°C	10% by volume of H ₂ in N ₂
B	4 h	450°C	10% by volume of H ₂ in N ₂
C	4 h	500°C	10% by volume of H ₂ in N ₂

- 5 The reduction was carried out using a gas mixture consisting of 10% by volume of H₂ in N₂. Here, the gas was passed at a temperature of about 400°C (batch A), 450°C (batch B) or 500°C (batch C) through the pellets for 4 hours in each case. The reduction was carried out under atmospheric pressure at a flow of 40 l/h.

The catalyst comprised: 7 g/l Au; 16 g/l; <1.5 g/l Cl; 10 g/l Pd.

15 Example 4

1.06 g of palladium acetate (224.49 g/mol), 0.66 g of gold acetate (374.10 g/mol) and 2.0 g of potassium acetate were dissolved in 15 ml of glacial acetic acid.

- 20 50 ml of TiO₂ supports (P25 pellets, DEGUSSA) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at 60°C on a rotary evaporator and residues of solvent were then removed at 60°C for 14 hours in a vacuum drying oven.

The reduction was carried out at 500°C for 1 hour using a gas mixture consisting of 10% by volume of H₂ in N₂, using a method similar to Example 1.

30

The catalyst contained: 7 g/l Au; 16 g/l K; <1 g/l Cl; 10 g/l Pd.

Example 5

1.06 g of palladium acetate (224.49 g/mol), 0.66 g of gold acetate (374.10 g/mol) and 4.0 g of potassium acetate were dissolved in 30 ml of glacial acetic acid at 60°C. 100 ml of TiO₂ supports (P25 pellets, DEGUSSA) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at 60°C on a rotary evaporator and residues of solvent were then removed at 60°C for 4 hours using an oil pump.

The reduction was carried out at 500°C for 1 hour using a gas mixture consisting of 10% by volume of H₂ in N₂, using a method similar to Example 1.

The catalyst contained: 3.5 g/l Au; 16 g/l K; <0.5 g/l Cl; 5 g/l Pd.

Comparative example 1

1.82 g of Na₂PdCl₄ (294.19 g/mol) and 0.64 g of NaAuCl₄ (361.76 g/mol) were dissolved in 32 ml of demineralized water. This solution was applied completely to 100 ml of SiO₂ supports (KA160 pellets, Süd-Chemie) with gentle agitation. To form a noble metal shell, the pretreated support was placed in a solution of 0.85 g of sodium hydroxide, NaOH, in 32 ml of demineralized water. The reaction mixture was left to stand overnight and the pellets were then washed free of chloride using demineralized water.

The catalyst was then reduced with an ethylene/nitrogen mixture (5% of ethylene in nitrogen) at 150°C for 5 hours.

To load the pellets with potassium ions, they were treated with a solution of 4 g of potassium acetate in 30 ml of water and the finished catalyst was dried for 2 hours in a rapid drier.

The catalyst comprised: 3.5 g/l Au; 16 g/l K; <0.5 g/l Cl; 6.6 g/l Pd.

Comparative example 2

1.06 g of palladium acetate (224.49 g/mol), 0.7 g of gold acetate (374.10 g/mol) and 4.0 g of potassium acetate were dissolved in 30 ml of glacial acetic acid at 60°C. 100 ml of TiO₂ supports (P25 pellets, DEGUSSA) were added to this solution. Subsequently, the major part of the acetic acid was firstly distilled off at 60°C on a rotary evaporator and residues of solvent were then removed at 60°C over a period of 4 hours using an oil pump.

Reduction was carried out using a gas mixture consisting of 10% by volume of H₂ and 90% by volume of N₂, in each case based on the volume, at 170°C for 1 hour (atmospheric pressure, gas flow: 40 l/h).

The catalyst comprised: 3.5 g/l Au; 16 g/l K; <0.5 g/l Cl; 5 g/l Pd.

Comparative example 3

0.53 g of palladium acetate (224.49 g/mol), 0.33 g of gold acetate (374.10 g/mol) and 2.0 g of potassium acetate were dissolved in 30 ml of glacial acetic acid. 50 ml of TiO₂ supports (XT25376 pellets, Norton) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at 70°C on a rotary evaporator, residues of solvent were then removed at 60°C with the aid of an oil pump and finally in a vacuum drying oven for a period of 14 hours, likewise at 60°C.

The reduction was carried out thermally without reducing gas (autoreduction). Here, nitrogen as flushing gas was passed through the pellets for 1 hour

at a temperature of about 500°C (atmospheric pressure, 40 l/h).

The catalyst comprised: 3.5 g/l Au; 16 g/l K; <0.5 g/l Cl; 5 g/l Pd.

Comparative example 4

1.06 g of palladium acetate (224.49 g/mol), 0.7 g of gold acetate (374.10 g/mol) and 4.0 g of potassium acetate were dissolved in 80 ml of glacial acetic acid at 60°C. 100 ml of SiO₂ supports (Aerosil 200 pellets, DEGUSSA) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at 60°C on a rotary evaporator and residues of solvent were then removed at 60°C over a period of 4 hours using an oil pump.

The reduction was carried out using a gas mixture consisting of 10% by volume of H₂ in N₂ for 1 hour at 500°C, using a method similar to Example 1.

The catalyst comprised: 3.5 g/l Au; 16 g/l K; <0.5 g/l Cl; 5 g/l Pd.

25

Comparative example 5

1.06 g of palladium acetate (224.49 g/mol), 0.7 g of gold acetate (374.10 g/mol) and 4.0 g of potassium acetate were dissolved in 80 ml of glacial acetic acid at 60°C. 100 ml of SiO₂ supports (KA160 pellets, Süd-Chemie) were added to this solution. Subsequently, the major part of the glacial acetic acid was firstly distilled off at 60°C on a rotary evaporator and residues of solvent were then removed at 60°C over a period of 4 hours using an oil pump.

The reduction was carried out using a gas mixture consisting of 10% by volume of H_2 in N_2 for 1 hour at 500°C, using a method similar to Example 1.

- 5 The catalyst comprised: 3.5 g/l Au; 16 g/l K; <0.5 g/l Cl; 5 g/l Pd.

Reactor tests for the gas-phase oxidation of ethylene and acetic acid to vinyl acetate:

10

The catalysts produced in the examples and comparative examples are tested in a fixed-bed tube reactor having a tube diameter of 2 cm. The reactor is heated externally by means of oil jacket heating. The reactor is typically charged with 15 ml of the shaped catalyst bodies. The reactor volume upstream and downstream of the catalyst bed is filled with glass spheres to reduce the dead volume. The gases are metered in via mass flow regulators for gases; the acetic acid is metered in using a mass flow regulator/vaporizer unit. The gases and the acetic acid are mixed in a packed gas mixing tube. The test apparatus is operated continuously.

20

25

The reaction is monitored continually by means of a gas chromatograph.

30

When a steady-state reaction has been achieved, i.e. the reactor temperature is constant and the concentrations of vinyl acetate and CO_2 in the product gas stream are constant, the recording of data is commenced.

35

The experiments were carried out using a reaction temperature in the range 150-170°C and a reaction pressure of 8-9 bar. The composition of the starting material stream was typically 60-80% by volume of ethylene, 10-20% by volume of N_2 , 10-20% by volume of acetic acid and 2-10% by volume of O_2 . A complete analysis of the output from the reactor was carried out

directly at the reactor outlet by means of on-line GC (2 column arrangement) and on-line IR.

The GC data were used to calculate the vinyl acetate selectivities S (= mol of VAM/(mol of VAM + 0.5* mol of CO_X , $X = 1$ or 2) and STY (space-time yield = g of VAM/l of cat. *h). Table 1 shows the results of the individual experiments in which the catalysts prepared in Examples 1 to 5 were tested.

10

Table 1: Testing of the catalysts in the synthesis of vinyl acetate

Example No.	T (°C)	p (bar)	O ₂ conc. (%)	S %	STY g/l*h
1A	170	9	5.2	96	1000
	160	9	5.2	98	1050
	155	9	5.2	98	1000
1B	170	9	5.2	97	700
	160	9	5.2	98	1150
1C	170	9	5.2	98	1300
2	170	9	5.2	96	1200
3A	160	9	5.2	89	1400
	150	9	5.2	98	1400
3B	160	9	5.2	95	1260
3C	160	9	5.2	96	1210
4	150	9	5.2	96	1100
5	160	9	5.2	95	940
Comp.1	170	9	5.1	88	850
Comp.2	160	9	5.2	80	870
Comp.3	170	9	5.2	77	<50
Comp.4	167	9	5.2	89	190
Comp.5	170	9	5.2	83	340

15 It can be seen from the table that the HTR treatment of catalysts in which a reducible support material is present significantly improves the selectivity and

yield in the vinyl acetate synthesis compared to the catalysts known from the prior art.

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Claims

1. A catalyst comprising palladium, at least one
alkali metal compound and, if desired, one or more
promoters on a porous support, obtainable by
loading the porous support, which comprises a
reducible metal oxide of elements of groups IIb,
IVb, Vb, VIb of the Periodic Table of the Elements
or ZnO or comprises a mixture of these oxides or a
mixed oxide of these elements in which zinc may
also be present, with at least one palladium
compound, subsequently carrying out a reduction at
a temperature of 300-600°C and additionally
applying at least one alkali metal compound and,
if desired, one or more promoters before or after
the reduction.
2. A catalyst as claimed in claim 1 which comprises
at least one potassium compound.
3. A catalyst as claimed in claim 1 or 2 which
additionally comprises Au, Ba and/or Cd and/or
their compounds as promoters.
4. A catalyst as claimed in one or more of claims 1
to 3, wherein the reducible support is TiO_2 .
5. A catalyst as claimed in one or more of claims 1
to 4, wherein the reduction is carried out for a
time in the range from 1 minute to 24 hours.
6. A catalyst as claimed in one or more of claims 1
to 5, wherein the reduction is carried out using
gaseous or vaporizable reducing agents.
7. A catalyst as claimed in one or more of claims 1
to 6, wherein the reducing agent for the reduction

is selected from the group consisting of H₂, CO, ethylene, NH₃, formaldehyde, methanol, hydrocarbons and their mixtures and mixtures of these reducing agents with inert gases.

- 5
8. A process for producing catalysts, which comprises loading the porous support, which comprises a reducible metal oxide of elements of groups IIIB, IVb, Vb, VIb of the Periodic Table of the Elements or ZnO or comprises a mixture of these oxides or a mixed oxide of these elements in which zinc may also be present, with at least one palladium compound, subsequently carrying out a reduction at a temperature of 300-600°C and additionally applying at least one alkali metal compound and, if desired, one or more promoters before or after the reduction.
- 10
9. The process as claimed in claim 8, wherein the catalyst comprises at least one potassium compound.
- 15
10. The process as claimed in claim 8 or 9 wherein the catalyst additionally comprises Au, Ba and/or Cd and/or their compounds as promoters.
- 20
11. The process as claimed in one or more of claims 8 to 10, wherein the reducible support is TiO₂.
- 25
12. The process as claimed in one or more of claims 8 to 11, wherein the reduction is carried out for a time in the range from 1 minute to 24 hours.
- 30
13. The process as claimed in one or more of claims 8 to 12, wherein the reduction is carried out using gaseous or vaporizable reducing agents.
- 35

14. The process as claimed in one or more of claims 8 to 13, wherein the reducing agent for the reduction is selected from the group consisting of H₂, CO, ethylene, NH₃, formaldehyde, methanol, hydrocarbons and their mixtures and mixtures of these reducing agents with inert gases.

15. The use of a catalyst obtainable as claimed in any of claims 8-14 for the preparation of vinyl acetate in the gas phase from ethylene, acetic acid or oxygen or oxygen-containing gases.

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DECLARATION

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Supplemental Sheet

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DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet

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Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor			
Given Name	Uwe	Middle Initial		Family Name	DINGERDISSEN	Suffix	
Inventor's Signature					Date		
Residence: City	Seeheim-Jugenheim	State		Country	Germany	Citizenship	DE
Post Office Address Linde-Weg 1, D-64342 Seeheim-Jugenheim, Germany							
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City	Seeheim-Jugenheim	State		Zip	D-64342	Country	Germany
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor			
Given Name	Klaus	Middle Initial		Family Name	KUHLEIN	Suffix	
Inventor's Signature	<i>Klaus Füll</i>				Date	1.9.07	
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Post Office Address							
City	Kelkheim	State		Zip	D-65779	Country	Germany
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor			
Given Name	Günter	Middle Initial		Family Name	DAMBECK	Suffix	
Inventor's Signature					Date		
Residence: City	Selters	State		Country	Germany	Citizenship	DE
Post Office Address Sackgasse 9, D-65618 Selters, Germany							
Post Office Address							
City	Selters	State		Zip	D-65618	Country	Germany
<input checked="" type="checkbox"/> Additional inventors are being named on supplemental sheet(s) attached hereto							

DECLARATION

ADDITIONAL INVENTOR(S)
Supplemental Sheet

Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor			
Given Name	Middle Initial	Family Name	Suffix e.g. Jr.				
Herhardt		GEISS					
Inventor's Signature	Gehard Geiss			Date	03.09.2001		
Residence: City	State	Country	Citizenship				
Liederbach		Germany	DE				
Post Office Address An der Untermühle 2, D-65835 Liederbach, Germany							
Post Office Address							
City	State	Zip	Country				
Liederbach		65835	Germany				
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor			
Given Name	Middle Initial	Family Name	Suffix e.g. Jr.				
Andrea		RUTSCH					
Inventor's Signature	Andrea Rutsch			Date	03.09.2001		
Residence: City	State	Country	Citizenship				
Mörfelden		Germany	DE				
Post Office Address Bürgermeister-Klingler-Strasse 5, D-64546, Mörfelden, Germany							
Post Office Address							
City	State	Zip	Country				
Mörfelden		D-64546	Germany				
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor			
Given Name	Middle Initial	Family Name	Suffix e.g. Jr.				
Stephan		WEIDLICH					
Inventor's Signature	Stephan Weidlich			Date	03.09.2001		
Residence: City	State	Country	Citizenship				
Frankfurt		Germany	DE				
Post Office Address Dürkheimer Strasse 37, D-65934, Frankfurt, Germany							
Post Office Address							
City	State	Zip	Country				
Frankfurt		D-65934	Germany				
Name of Additional Joint Inventor, if any:				<input type="checkbox"/> A petition has been filed for this unsigned inventor			
Given Name	Middle Initial	Family Name	Suffix e.g. Jr.				
Inventor's Signature				Date			
Residence: City	State	Country	Citizenship				
Post Office Address							
Post Office Address							
City	State	Zip	Country				
<input type="checkbox"/> Additional inventors are being named on supplemental sheet(s) attached hereto							

DECLARATION FOR
UTILITY OR DESIGN
- PATENT APPLICATION

Attorney Docket Number	RUH-284
First Named Inventor	HAGEMEYER et al
COMPLETE IF KNOWN	
Application Number	PCT/EP00/02455
Filing Date	March 21, 2001
Group Art Unit	
Examiner Name	

☒ Declaration Submitted with Initial Filing OR ☐ Declaration Submitted after Initial Filing

As a below named inventor, I hereby declare that:

My residence, past office address, and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

CATALYSTS FOR THE GAS-PHASE OXIDATION OF ETHYLENE AND ACETIC ACID TO VINYL ACETATE, A PROCESS FOR PRODUCING THEM AND THEIR USE

(Date of the invention)

the specification of which

☐ is attached hereto
OR☒ was filed on (MM/DD/YYYY) 03/21/01 as United States Application Number or PCT International

Application Number PCT/EP00/02455 and was amended on (MM/DD/YYYY) (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37 Code of Federal Regulations, § 1.56.

I hereby claim foreign priority benefits under Title 35, United States Code § 119 (a)-(e) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365 (a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached?	
199 14. 066.9	Germany	3/27/99	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	YES <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>	NO <input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>

☐ Additional foreign application numbers are listed on a supplemental priority sheet attached hereto:

I hereby claim the benefit under Title 35, United States Code § 119(e) of any United States provisional application(s) listed below.

Application Number(s)	Filing Date (MM/DD/YYYY)	<input type="checkbox"/> Additional provisional application numbers are listed on a supplemental priority sheet attached hereto.

(Page 1 of 5)

Burden Hour Statement: This form is estimated to take 0.4 hours to complete. Time will vary depending upon the needs of the individual case. Any comments on the amount of time you are required to complete this form should be sent to the Chief Information Officer, Patent and Trademark Office, Washington, DC 20231. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS SEND TO Commissioner of Patents and Trademarks, Washington, DC 20231



Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

DECLARATION

I hereby claim the benefit under Title 35, United States Code §120 of any United States application(s), or §365(c) of any PCT international application designating the United States of America, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT international application in the manner provided by the first paragraph of Title 35, United States Code §112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application.

U.S. Parent Application Number	PCT Parent Number	Parent Filing Date (MM/DD/YYYY)	Parent Patent Number (if applicable)

☐ Additional U.S. or PCT international application numbers are listed on a supplemental priority sheet attached hereto.

As a named inventor, I hereby appoint the following registered practitioner(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:

Name	Registration Number	Name	Registration Number
Bierman, Muserlian and Lucas	18,818	④	
Jordan B. Bierman	18,629		
Charles A. Muserlian	19,683		
Donald C. Lucas	31,275		

☐ Additional registered practitioner(s) named on a supplemental sheet attached hereto.

Direct all correspondence to:

Name	Charles A. Muserlian		
Address	Bierman, Muserlian and Lucas		
Address	600 Third Avenue		
City	New York	State	NY
Country	U.S.A.	Telephone	212-661-8000
		Fax	212-661-8002

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true, and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Name of Sole or First Inventor:

☐ A petition has been filed for this unsigned inventor

Given Name	Middle Initial	Family Name	Suffix e.g. Jr.
Alfred		HAGEMEYER	

Inventor's Signature	Alfred Hagemeyer	Date	09/04/01
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Residence: City	Sunnyvale	State	CA	Country	US	Citizenship	US
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City	Sunnyvale	State		Zip	94086	Country	US
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☒ Additional inventors are being named on supplemental sheet(s) attached hereto